

الطباطبائی

GenCore version 5.1.4_P5-1578
 Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - protein search, using sw model

Run on: April 8, 2003, 14:23:18 ; Search time 40 Seconds
 (without alignments)
 1815,540 Million cell updates/sec

Title: US-09-001-737-8
 Perfect score: 2663
 Sequence: I MAKEIKFSADARRAMVRGVDF.....TPAPAMPAGMDPGHMGGG 545

Scoring table: BLOSUM62
 Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0
 Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
 Maximum Match 100%
 Listing first 45 summaries

Database : A_GeneSeq_101002-*
 1: /\$IDS2/gcdata/geneseq/geneseqp-embl/RA1980.DAT: *
 2: /\$IDS2/gcdata/geneseq/geneseqp-embl/RA1981.DAT: *
 3: /\$IDS2/gcdata/geneseq/geneseqp-embl/RA1982.DAT: *
 4: /\$IDS2/gcdata/geneseq/geneseqp-embl/RA1983.DAT: *
 5: /\$IDS2/gcdata/geneseq/geneseqp-embl/RA1984.DAT: *
 6: /\$IDS2/gcdata/geneseq/geneseqp-embl/RA1985.DAT: *

11 1880.5 70.6 539 22 AAGB81848
 12 1847.5 69.4 539 20 AAY21916
 13 1833.5 68.9 540 23 ABB3910
 14 1819 68.3 539 20 AAY21906
 15 1702 63.9 541 20 AAY21910
 16 1691.5 63.5 548 18 AAW16678
 17 1671 62.7 541 20 AAY1909
 18 1671 62.7 541 23 ABB7515
 19 1654 62.1 545 20 AAY123915
 20 1653.5 62.1 540 18 AAW31100
 21 1652.5 62.1 948 22 AAB31611
 22 1652 62.0 544 21 AAY5747
 23 1647.5 61.9 540 9 AAB1135
 24 1647.5 61.9 540 16 AAB81610
 25 1647.5 61.9 540 19 AAW44702
 26 1647.5 61.9 540 20 AAY23911
 27 1647.5 61.9 540 21 AAY31332
 28 1647.5 61.9 540 22 AAB1175
 29 1647.5 61.9 540 22 AAB8118
 30 1647.5 61.9 540 22 AAB31606
 31 1647.5 61.9 540 23 AAM0750
 32 1647.5 61.9 560 9 AAB80215
 33 1647.5 61.9 560 22 AAB1169
 34 1647.5 61.9 608 22 AAB11614
 35 1643 61.7 528 19 AAW0144
 36 1643 61.7 528 20 AAY14891
 37 1643 61.7 523 23 ABB73497
 38 1643 61.7 639 21 AAB01790
 39 1642.5 61.7 639 22 AAB11609
 40 1641 61.6 505 20 AAY23901
 41 1640 61.6 508 19 AAW51070
 42 1640 61.6 508 20 AAW51001
 43 1640 61.6 548 22 AAB50536
 44 1639.5 61.6 500 23 AAW16511
 45 1639.5 61.6 500 23 AAW16194
 46 1639.5 61.6 500 16 AAB7384

S.	epidermidis	ope
Amino acid sequenc		
Staphylococcus epi		
Amino acid sequenc		
Amino acid sequenc		
Lawsonia intracellularis		
Amino acid sequenc		
M. vaccae	GroEL hom	
Amino acid sequenc		
Mycobacterium sp. h		
Amino acid sequenc		
Neisseria meningitidis		
Sequence of Mycoba		
Mycobacterium tube		
Amino acid sequenc		
Amino acid sequenc		
Mycobacterium tube		
Amino acid sequenc		
Mycobacterium tube		
Amino acid sequenc		
Mycobacterium tube		
Sequence of Mycoba		
Amino acid sequenc		
M. vaccae antigen		
Amino acid sequenc		
M. vaccae	GroEL hom	
Heat shock protein		
Streptococcus pneumoniae		
M. leprae	GroEL 9	

```

10: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1988.DAT:*
11: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1989.DAT:*
12: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1990.DAT:*
13: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1991.DAT:*
14: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1992.DAT:*
15: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1993.DAT:*
16: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1994.DAT:*
17: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1995.DAT:*
18: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1996.DAT:*
19: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1997.DAT:*
20: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1998.DAT:*
21: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa1999.DAT:*
22: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa2000.DAT:*
23: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa2001.DAT:*
24: /SDS2/gcgdata/geneseq/geneseq-emb1/Aa2002.DAT:*

```


CC Streptococcus bacteria, particularly *S. agalactiae* and *S. pyogenes*.
 CC Nucleic acids encoding (I) are used to detect Streptococcus in a
 CC biological sample. (I) is used to determine whether a compound binds to
 CC (I). A composition comprising (I) or a nucleic acid encoding (I), may be
 CC used as a vaccine or diagnostic composition. The disease caused by
 Streptococcus that is prevented or treated may be meningitis. Nucleic
 acid encoding (I) may be used to recombinantly produce (I) and may be
 CC used in gene therapy. Antibodies to (I) are used for affinity
 chromatography, immunoassays, and distinguishing/identifying
 CC Streptococcus proteins.

QY 241 TNRPLLIADDVGEALPFLVNLKIRGTFNVAVRAKGPGFDRKAMLEDAITLGCVIT 300
 Db 243 TNRPLLIADDVGEALPFLVNLKIRGTFNVAVRAKGPGFDRKAMLEDAITLGCVIT 302

QY 301 EDIGLEKIDAMTALGQAKTIVDSDTIVGEGSSSEALANIAILKSOLETTSDFDR 360
 Db 303 EDIGLEKIDAMTALGQAKTIVDSDTIVGEGSSSEALANIAILKSOLETTSDFDR 362

QY 361 EKQERLAKLAGGVAVIKVAGPTEALKEMKIRIEDALANATRAVERGTVAGGTALITY 420
 Db 363 EKQERLAKLAGGVAVIKVAGPTEALKEMKIRIEDALANATRAVERGTVAGGTALITY 422

QY 421 IEKVALELEGDDATGRNIVRALLEPVRQALNAGIEGSVTDKLNNSPAGGFNATG 480
 Db 423 IEKVALELEGDDATGRNIVRALLEPVRQALNAGIEGSVTDKLNNSPAGGFNATG 482

QY 481 EWMVKGKIDDPVYVTSALONAASVSLITTEAVWANKPEPATPAPAMPAGMDGMM 580
 Db 483 EWMVKGKIDDPVYVTSALONAASVSLITTEAVWANKPEPATPAPAMPAGMDGMM 542

QY 541 GG 542

Db 543 GG 544

RESULT 3

ABP28528 ABP28528 standard; Protein; 540 AA.
 XX
 AC ABP28528;
 DT 02-JUL-2002 (first entry)
 DE Streptococcus polypeptide SEQ ID NO 6232.
 XX
 KW Streptococcus; GAS; GBS; group B streptococcus; Streptococcus agalactiae;
 KW group A streptococcus; Streptococcus pyogenes; antibacterial;
 KW antiinflammatory; infection; vaccine; meningitis; gene therapy.
 XX
 OS Streptococcus agalactiae.
 XX
 PN WO200234771-A2.
 PD 02-MAY-2002.
 XX
 PF 29-OCT-2001; 2001WO-GB04789.
 XX
 PR 27-OCT-2000; 2000GB-0026133.
 PR 24-NOV-2000; 2000GB-002727.
 PR 07-MAR-2001; 2001GB-0005640.
 XX
 PA (CHIR-) CHIRON SPA.
 PA (GENO-) INST GENOMIC RES.
 XX
 PI Telford J, Maignani V, Margarit Ros YI, Grandi G, Fraser C;
 PI Tellelin H;
 XX
 DR WPI; 2002-352536/38.
 DR N-PSDB; ABN69159.
 XX
 PT New Streptococcus protein for the treatment or prevention of infection
 PT or disease caused by Streptococcus bacteria, such as meningitis, and
 PT for detecting a compound that binds to the protein -
 XX
 PS Claim 1; Page 3795; 4525pp; English.

RESULT 4

AY23902 AY23902 standard; Protein; 541 AA.
 XX
 AC AY23902;
 DT 22-SEP-1999 (first entry)
 XX
 DE Streptococcus pneumoniae heat shock protein (Hsp)60-2.
 XX
 KW Heat shock protein; Hsp60-2; immune response; immunological carrier;
 KW cancer control; tumour; sarcoma; cancer; gene therapy.
 XX
 OS Streptococcus pneumoniae.

CC The invention relates to a protein (ABP2413-ABP3089) from group B
 CC streptococcus/GBS (Streptococcus agalactiae) or group A streptococcus/GAS
 CC (Streptococcus pyogenes), comprising one of 543 sequences (S1), given in
 CC the specification. The proteins have antibacterial and antiinflammatory
 CC activity. (I), nucleic acids encoding (I), ABN6044 ABN7152 and
 CC antibodies that bind (I) are used in the manufacture of medicaments for
 the treatment or prevention of infection or disease caused by

PN WO935270-A1.
 XX
 PD 15-JUL-1999.
 XX
 XX
 PR 29-DEC-1998; 98WO-CA01203.
 XX
 PR 31-DEC-1997; 97US-0001737.
 XX
 PA (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.
 XX
 PT Mizzen L, Wisniewski J;
 XX
 WPI; 1999-430397/36;
 DR N-PSDB; AAX86153.
 XX
 New nucleic acid encoding heat shock protein-60 from *Streptococcus*, useful in vaccines, as carriers for other immunogens, as anticancer agents and for diagnosis
 XX
 PS Claim 11; FIG 2A-B; 176pp; English.
 CC The present sequence represents a heat shock protein, designated Hsp60-2. The protein, its fragments, variants and fusion proteins, are used to elicit or enhance an immune response against *Streptococcus*, and to elicit a similar response to a target antigen fused to the protein. Unlike other immunological carriers, Hsp60 proteins are not immunosuppressive so provide an increased response to any conjugated or fused antigen. Also, where used for cancer control, they lack the side effects associated with endotoxins. They can also be used to detect specific antibodies and in treatment or prevention of tumours (e.g. sarcoma or cancers of breast, ovary, prostate, lung, pancreas or liver). The Hsp60 polynucleotide is used for recombinant production of the protein, as a source of primers and probes for detecting *Streptococci* in standard hybridization/amplication assays, and therapeutically in gene therapy vectors.
 XX
 SQ Sequence 541 AA:
 Query Match 88.5%; Score 2358; DB 20; Length 541;
 Best Local Similarity 87.9%; Pred. No. 3.1e-144; Mismatches 479; Indels 4; Gaps 1;
 Matches 129; Conservative 33; MisMatches 29; Indels 4; Gaps 1;
 PT 1 MAKEIKFSDARABAMVRGUDMLADIVKVTGPKGRNVVLEKAFGSPPLITNDGYTIKEIE 60
 1 MAKEIKFSDARABAMVRGUDMLADIVKVTGPKGRNVVLEKAFGSPPLITNDGYTIKEIE 60
 XX
 QY 61 LEFHFNENGGAKVSEVASKNDIAGGGTTATVLQIVIGLNVNAGANPIGKIGRIGIE 120
 61 LEFHFNENGGAKVSEVASKNDIAGGGTTATVLQIVIGLNVNAGANPIGKIGRIGIE 120
 Db 61 LEFHFNENGGAKVSEVASKNDIAGGGTTATVLQIVIGLNVNAGANPIGKIGRIGIE 120
 PT Screening for compounds that stimulate Th1-like responses in CD4+ T lymphocyte cells.
 XX
 PS Example 15; FIG 15A-B; 88pp; English.
 XX
 The present sequence represents a fusion protein comprising a *Streptococcus pneumoniae* heat shock protein (Hsp) 65 fused to a HPV16 E7 protein. The fusion protein is used in the method of the invention. The specification describes a method of determining whether a compound stimulates a Th1-like response. Th1 cells are a subset of CD4+ T lymphocyte cells. The method comprises contacting naive lymphocytes in vitro with a fusion protein comprising at least a fragment of Hsp, and then detecting the Th1-like response exhibited by the cell sample. The proteins which may be used in the method of the invention are Hsp65, Hsp40, Hsp60, and Hsp70. The method may be used to identify compounds that stimulate Th1-like responses in response to microbial pathogens.
 XX
 SQ Sequence 641 AA:
 Query Match 88.5%; Score 2358; DB 22; Length 641;
 Best Local Similarity 87.9%; Pred. No. 4.1e-144; Mismatches 479; Indels 4; Gaps 1;
 Matches 129; Conservative 33; MisMatches 29; Indels 4; Gaps 1;
 PT 1 MAKEIKFSDARABAMVRGUDMLADIVKVTGPKGRNVVLEKAFGSPPLITNDGYTIKEIE 60
 1 MAKEIKFSDARABAMVRGUDMLADIVKVTGPKGRNVVLEKAFGSPPLITNDGYTIKEIE 60
 XX
 QY 361 ELQBLRALKLGGVAVIKVQATTEBEKMKRLEDDLNTRAVVEGIVAGGGTALANV 420
 361 ELQBLRALKLGGVAVIKVQATTEBEKMKRLEDDLNTRAVVEGIVAGGGTALANV 420
 Db 421 IERVVALELEGGDGTNTYVRLAEEFVROTALNAGVEGSVVIKNSPAGTGPNAATG 480
 421 IERVVALELEGGDGTNTYVRLAEEFVROTALNAGVEGSVVIKNSPAGTGPNAATG 480
 DB 421 IERVVALELEGGDGTNTYVRLAEEFVROTALNAGVEGSVVIKNSPAGTGPNAATG 480
 XX
 PA (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.
 XX
 PT Siegel M, Chu NR, Mizzen LA;
 XX
 DR N-PSDB; AAF250156.
 XX
 PT 08-JUL-1999; 99US-0143757.
 XX
 PA (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.
 XX
 DR WPI; 2001-13831/4.
 XX
 PT Screening for compounds that stimulate Th1-like responses in CD4+ T lymphocyte cells.
 XX
 PS Example 15; FIG 15A-B; 88pp; English.
 XX
 The present sequence represents a fusion protein comprising a *Streptococcus pneumoniae* heat shock protein (Hsp) 65 fused to a HPV16 E7 protein. The fusion protein is used in the method of the invention. The specification describes a method of determining whether a compound stimulates a Th1-like response. Th1 cells are a subset of CD4+ T lymphocyte cells. The method comprises contacting naive lymphocytes in vitro with a fusion protein comprising at least a fragment of Hsp, and then detecting the Th1-like response exhibited by the cell sample. The proteins which may be used in the method of the invention are Hsp65, Hsp40, Hsp60, and Hsp70. The method may be used to identify compounds that stimulate Th1-like responses in response to microbial pathogens.
 XX
 SQ Sequence 641 AA:
 Query Match 88.5%; Score 2358; DB 22; Length 641;
 Best Local Similarity 87.9%; Pred. No. 4.1e-144; Mismatches 479; Indels 4; Gaps 1;
 Matches 129; Conservative 33; MisMatches 29; Indels 4; Gaps 1;
 PT 1 MAKEIKFSDARABAMVRGUDMLADIVKVTGPKGRNVVLEKAFGSPPLITNDGYTIKEIE 60
 1 MAKEIKFSDARABAMVRGUDMLADIVKVTGPKGRNVVLEKAFGSPPLITNDGYTIKEIE 60
 XX
 QY 61 LEFHFNENGGAKVSEVASKNDIAGGGTTATVLQIVIGLNVNAGANPIGKIGRIGIE 120
 61 LEFHFNENGGAKVSEVASKNDIAGGGTTATVLQIVIGLNVNAGANPIGKIGRIGIE 120

The present invention relates to nucleic acids (AAM0102-1-AAM0114), which are essential for the encoding of polypeptides (AAM0102-AAM0114), which are essential for the viability of a bacterial cell wall. The acronym CEG stands for "Conserved Essential Gene". The nucleic acids are useful for detecting the presence of proteins essential for the viability of a bacterial cell wall in samples such as cells, tissues, biological fluids, blood, serum, nose, ear or throat swabs with ligands, and for detecting corresponding target nucleic acid molecules with complementary sequences. The nucleic acids are also useful for determining whether a genomic nucleotide sequence of interest is essential for viability of bacterial cell or whether it resides within an operon, by integrating an exogenous nucleotide sequence comprising a portion of an open reading frame of the genomic sequence of interest (comprising 200-500 base pairs) into the genomic sequence of interest which confers a selectable phenotype to the cell, and determining cell viability with a selection agent such as chloramphenicol. The nucleic acids and proteins are also useful as vaccines and for treating bacterial infections with gene therapy and antisense therapy. The nucleic acids also enable identification of targets suitable for the treatment of

QY	121	TATATPVALAKIAQPVSGKEIAQAVAYSSRSERKVEYSEAMERYGNDGVTIESRG 180	CC
Db	121	TAVAATVEALKNNIVPVANKELIAQAVAYSSERKVEYSEAMERYGNDGVTIESRG 180	CC
QY	181	METLEVEWGGQFDRGYLSQYMTDNEKADLENPILTDKVNQDQPLPLEELK 240	CC
Db	181	METEVEWGGQFDRGYLSQYMTDNEKADLENPILTDKVNQDQPLPLEELK 240	CC
Db	181	METEVEWGGQFDRGYLSQYMTDNEKADLENPILTDKVNQDQPLPLEELK 240	CC
QY	241	TNPPLIADDQGEALPVLINKRGFNVAVKAGFGRKAMLEDTALGTVIT 300	CC
Db	241	TNPPLIADDQGEALPVLINKRGFNVAVKAGFGRKAMLEDTALGTVIT 300	CC
Db	241	TNPPLIADDQGEALPVLINKRGFNVAVKAGFGRKAMLEDTALGTVIT 300	CC
QY	301	EDGLELKDMATLGQAKITVDKSTVWEGGSSEATANRALKSOLETTSDFR 360	CC
Db	301	EDGLELKDMATLGQAKITVDKSTVWEGGSSEATANRALKSOLETTSDFR 360	CC
Db	301	EDGLELKDMATLGQAKITVDKSTVWEGGSSEATANRALKSOLETTSDFR 360	CC
QY	361	EKQERLAKLAGVAVTGVGAPTEALEMKURLEDTANTRAVEGIVGGTALV 420	CC
Db	361	EKQERLAKLAGVAVTGVGAPTEALEMKURLEDTANTRAVEGIVGGTALV 420	CC
Db	361	EKQERLAKLAGVAVTGVGAPTEALEMKURLEDTANTRAVEGIVGGTALV 420	CC
QY	421	IEKVALELEGDTGRNIVRALERPEVQTLNAGYEGSVWIDKLNKPAGTGAATG 480	CC
Db	421	IEKVALELEGDTGRNIVRALERPEVQTLNAGYEGSVWIDKLNKPAGTGAATG 480	CC
QY	481	EDMDTIGTIDIPKVRSALQNAASVSLITTEAVVNPAPPAPAPMPAGMAGM 540	CC
Db	481	EDMDTIGTIDIPKVRSALQNAASVSLITTEAVVNPAPPAPAPMPAGMAGM 540	CC
Qy	541	GGMGG 545	CC
Db	537	GGMGG 541	CC
RESULT 6			
AM01101			
ID	AM01101 standard; Protein: 540 AA.		
XX			
AC	AM01101;		
XX			
DE	02-OCT-2001 (first entry)		
XX			
CFE	104 protein sequence.		
XX			
KW	Antibacterial; vaccine; gene therapy; bacterial cell wall viability; antisense therapy; antibiotic resistance.		
XX			
OS	Streptococcus pneumoniae.		
XX			
WO20019721-A2.			
XX			
PD	12-JUL-2001.		
XX			
PF	29-DEC-2000; 2000MO-US35604.		
XX			
PR	30-DEC-1999; 99US-0174089.		
XX			
PA	(BRIM) BRISTOL-MYERS SQUIBB CO.		
XX			
PT	Dougherty TJ, Pucci MJ, Dougherty BA, Davison DB, Brucolieri RE; Thanassi JA;		
PT	WPI: 2001+96721/54.		
DR	N-PSDB; AAH90800.		
XX			
PT	Nucleic acids encoding conserved essential genes involved in bacterial replication which are potential targets for the treatment of antibiotic resistant bacterial infections.		
PS	Claim 27; Pages 356-358; 380pp; English.		
AC	ABR53701.		
RESULT 7			
PT	ABBS3701		
ID	ABBS3701 standard; Protein: 542 AA.		
XX			

RESULT 10	AAV23930	ID AAV23930	DE Consensus mino:acid sequence of a heat shock protein.
XX	AC AAY23930:	XX	XX
DT 22-SEP-1999	(first entry)	DE	Heat shock protein; Hsp; immune response; immunological carrier; cancer control; tumour; sarcoma; cancer; gene therapy.
XX	OS Synthetic.	XX	OS
PN W0935370-A1.	XX	PN	XX
PD 15-JUL-1999.	XX	PD	XX
PF 29-DEC-1998:	98W0-C0A01203.	PF	XX
PR 31-DEC-1997:	97US-0001737.	PR	XX
XX	(STRE-) STRESSGEN BIOTECHNOLOGIES CORP.	XX	XX
PI Mizzen L, Wisniewski J;	XX	PI	XX
DR WPI: 1999-430397/36.	XX	DR	XX
XX	PT New nucleic acid encoding heat shock protein-60 from <i>Streptococcus</i> , useful in vaccines, as carriers for other immunogens, as anticancer agents and for diagnosis	PT	XX
CC	PS Disclosure: FIG 10A-B: 17pp; English.	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
CC	CC	CC	XX
SQ Sequence 545 AM:	XX	SQ	XX
Query Match 71.5%; Score 1904.5; DB 20; Length 545; Matches 392; Conservative 6; Mismatches 80; Indels 9; Gaps 6;	XX	Query Match 71.5%; Score 1904.5; DB 20; Length 545; Matches 392; Conservative 6; Mismatches 80; Indels 9; Gaps 6;	XX
QY 1 MAKEIKFSADAKANVYRGMDLADTVKVLGPKGRNVLKARGSPPLITNGVTAKEIE 60	XX	QY 1 MAKEIKFSADAKANVYRGMDLADTVKVLGPKGRNVLKARGSPPLITNGVTAKEIE 60	XX
Db 1 MAKDKEEERRMRMGRNMLADKVKGPKGRNVLKSGPAPITKIGVIVANIE 60	XX	Db 1 MAKDKEEERRMRMGRNMLADKVKGPKGRNVLKSGPAPITKIGVIVANIE 60	XX
QY 121 TATAATVAKLAIAQPVSGKEIAQAVYASSR-SERGVETISAMERVNGVITIERSR 179	XX	QY 121 TATAATVAKLAIAQPVSGKEIAQAVYASSR-SERGVETISAMERVNGVITIERSR 179	XX
Db 121 KAVDVAWEELKAIKAPVKEIKAQVATISANGDERRIGELIABAMKVGEGVITVEBGK 180	XX	Db 121 KAVDVAWEELKAIKAPVKEIKAQVATISANGDERRIGELIABAMKVGEGVITVEBGK 180	XX
QY 180 GMETELEVVEGMQFDGRGIVSQMWMNDNEKVNADLMPFLITDKKYSNTIDQIPLBEVL 239	XX	QY 180 GMETELEVVEGMQFDGRGIVSQMWMNDNEKVNADLMPFLITDKKYSNTIDQIPLBEVL 239	XX
QY 181 TLETELEVEMDQFDGRGIVSPFIDSEROKAELDPLILDKNSNIDQIPLBEVA 240	XX	QY 181 TLETELEVEMDQFDGRGIVSPFIDSEROKAELDPLILDKNSNIDQIPLBEVA 240	XX
PS	Sequence 545 AM:	PS	Sequence 545 AM:
DR WPI: 2001-15495/33.	XX	DR WPI: 2001-15495/33.	XX
DR N-PSDB; AAH52698.	XX	DR N-PSDB; AAH52698.	XX
PT Nucleic acids encoding polypeptides from <i>Staphylococcus epidermidis</i> , useful for vaccinating against infections, e.g. endocarditis -	XX	PT Nucleic acids encoding polypeptides from <i>Staphylococcus epidermidis</i> , useful for vaccinating against infections, e.g. endocarditis -	XX
PS Claim 18; Page 243; 2188pp; English.	XX	PS Claim 18; Page 243; 2188pp; English.	XX
XX	AAH52304 to AAH53970 represent nucleic acids (I) encoding polypeptides (II), given in AAH81154 to AAH81120, from <i>Staphylococcus epidermidis</i> , (I) and (II) can have antibacterial activity and therefore can be used in vaccination. The nucleic acids (I) may be used to produce the S. epidermidis polypeptides (II) via the production of vectors containing them which are used to produce host cells which express the polypeptides. The polypeptides (II) (and/or nucleic acid) may then be used to vaccinate subjects and to raise antibodies against the bacterium. The polypeptides may also be used to assay for other inhibitors of the activity and therefore identify compounds that may be used for the treatment of S. epidermidis infections, e.g. endocarditis. AAH53970 to AAH55030 represent specifically claimed S. epidermidis genomic DNA	XX	AAH52304 to AAH53970 represent nucleic acids (I) encoding polypeptides (II), given in AAH81154 to AAH81120, from <i>Staphylococcus epidermidis</i> , (I) and (II) can have antibacterial activity and therefore can be used in vaccination. The nucleic acids (I) may be used to produce the S. epidermidis polypeptides (II) via the production of vectors containing them which are used to produce host cells which express the polypeptides. The polypeptides (II) (and/or nucleic acid) may then be used to vaccinate subjects and to raise antibodies against the bacterium. The polypeptides may also be used to assay for other inhibitors of the activity and therefore identify compounds that may be used for the treatment of S. epidermidis infections, e.g. endocarditis. AAH53970 to AAH55030 represent specifically claimed S. epidermidis genomic DNA

CC Polynucleotide sequences from the present invention, AAH55091 to
 CC AAH55098 represent oligonucleotide sequences and primers which are used
 CC in the exemplification of the present invention.
 CC N.B. The present invention specifically claims all the polynucleotide
 CC sequences given in the sequence listing of the present specification,
 CC however the sequence listing only goes up to SEQ ID NO:4456, so even
 CC though sequences are given in the disclosure for SEQ ID NO:4465 to 4472,
 XX no sequences are present for SEQ ID NO:4455 to 4464.

SQ Sequence 539 AA;

Query Match 70.6%; Score 1880.5; DB 22; Length 539;
 Best Local Similarity 68.1%; Pred. No. 2.2e-113; PS
 Matches 368; Conservative 85; Mismatches 86; Indels 1; Gaps 1;

Qy 1 MAKEIKFSADARAAMVRGVMDLADPKVTLGKGRNVLEKARGSPPLITDGVTAKIEIE 60
 Db 1 MARDKLFSESDARQMLRQVDRKLANKAVKTVGPKGRNVVNDKDTPTPLITDGVTAKIEIE 60
 Qy 61 LEDHENMKAKLYSEVASKTDIAGDTTAVLQATVHEGLKNTAGANPPIGIRGIE 120
 Db 61 LEDPYENNGAKLYVEVANKTNEIAGDTTATVLAQSMIQEGLKNTSGANPVGRLRGID 120
 Qy 121 TATATAVEALKIAQPVSGKIAQAVAVSSPKERYGETISAMERYNGDVTIBSRG 180
 Db 121 KAVKVALEHNSOKYENKEIAGQVYISADEIGRVISSAMDRKVNGDVTIBESNG 180
 Qy 181 MEBLENEVGMDPDRGLSOMYMTNEKADLENPFLITOKVNSIDUDLLEVRK 240
 Db 181 ENTELEVEMQFDGQYQSPYMWTSKMDKMLABLERPYLTWTKKSFQDILPLBQVO 240
 Qy 241 TAPPLPLIADDVGEALPLTYLNUKRGFTNIVAVKARGFGRKAMLDIAITGGVIT 300
 Db 241 ASRPLIVADEVEGDAFLNTIVLNRMGRFTAVAKVAGPKGDRKAMLDIAITGGQIT 300
 Qy 301 EDIGLGLKDTATMPALGQAKATVKDSTVYEGSSSEIATNRLALKSOLETTSDFR 360
 Db 301 DQGELKLKDASDMLGATKWKVNEKDTVTDGNDENNIDARVGQKIAQIEPDSEFDK 360
 Qy 361 EKLOERLAKLAGVAVKVGAPTEALKEMKLRTDNLNTRAAVEGTVAGSGTALIV 420
 Db 361 EKLOERLAKLAGVAVKVGASTELKERKLRIEDALNSTRAVEGTVAGGTLVNI 420
 Qy 421 IEKVALELEGDDATGNTTIALELPYRQALNAGYGSVYDVKUNSPAGTGFNATG 480
 Db 421 YOKVSEKAGDVETGVNIVNLQKAPYQDIAENAGLEGSIIVERLKHAEAGVGFNATN 480
 Qy 481 ENWDMKTSITDIDVYKVRSLQNLQNSASVSLTTEAVVANKPEPATPAPAMPMDPGMM 540
 Db 481 ENVNMLEEGIVDPTKVRSALQHASYVAMFLTEAUVVASIPENNEBPGMGM-PGMM 539

RESULT 12

AY23916 31-DEC-1997; 97US-0001737;
 XX (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.
 XX Mizen L, Wisniewski J;
 XX DR WPI: 1999-430397/36.

PR 31-DEC-1997; 97US-0001737;
 XX PA (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.
 XX PT Mizzen L, Wisniewski J;
 XX DR WPI: 1999-430397/36.

XX New nucleic acid encoding heat shock protein-60 from *Streptococcus*,
 PR useful in vaccines, as carriers for other immunogens, as anticancer
 PR agents and for diagnosis

XX Disclosure: Fig 10A-E, 176pp;

XX English.

XX AAU23905-30 represent heat shock proteins (Hsps). The specification
 CC describes *Streptococcal* Hsps, designated Hsp60. These proteins, their
 fragments, Variants and fusion proteins, are used to elicit or enhance
 CC an immune response against *streptococcus*, and to elicit a similar
 CC response to a target antigen fused to the protein. Unlike other
 CC immunological carriers, Hsp60 proteins are not immunosuppressive so
 CC provide an increased response to any conjugated or fused antigen. Also,
 CC where used for cancer control, they lack the side effects associated
 CC with endotoxins. They can also be used to detect specific antibodies
 CC and in treatment or prevention of tumours (e.g. sarcoma, or cancers of
 CC breast, ovary, prostate, lung, pancreas or liver). The Hsp60
 CC polynucleotide is used for recombinant production of the protein, as
 CC a source of primers and probes for detecting *streptococci* in standard
 CC hybridization/amplification assays, and therapeutically in gene
 XX therapy vectors.

SQ Sequence 539 AA;

Query Match 69.4%; Score 1847.5; DB 20; Length 539;
 Best Local Similarity 68.3%; Pred. No. 2.9e-111; PS
 Matches 371; Conservative 78; Mismatches 87; Indels 7; Gaps 4;

Qy 1 MAKEIKFSADARAAMVRGVMDLADPKVTLGKGRNVLEKARGSPPLITDGVTAKIEIE 60
 Db 1 MVKQIKFSADARAAMVRGVMDLADPKVTLGKGRNVLEKARGSPPLITDGVTAKIEIE 60
 Qy 61 LEDHENMKAKLYSEVASKTDIAGDTTAVLQATVHEGLKNTAGANPPIGIRGIE 120
 Db 61 LEDPYENNGAKLYVEVANKTNEIAGDTTATVLAQSMIQEGLKNTSGANPVGRLRGID 120
 Qy 121 TATATAVEALKIAQPVSGKIAQAVAVSSPKERYGETISAMERYNGDVTIBSRG 180
 Db 121 KAVKVALEHNSOKYENKEIAGQVYISADEIGRVISSAMDRKVNGDVTIBESNG 180
 Qy 181 SRMTEEL - GMFDQYQSPYMWTSKMDKMLABLERPYLTWTKKSFQDILPLBQ 238
 Db 181 SRMTEEL - GMFDQYQSPYMWTSKMDKMLABLERPYLTWTKKSFQDILPLBQ 238
 Qy 238 VLKNTPLIADDVGEALPLTYLNUKRGFTNIVAVKARGFGRKAMLDIAITGGT 297
 Db 239 VVOSNRLIVADEVEGDAFLNTIVLNRMGRFTAVAKVAGPKGDRKAMLDIAITGAQ 298
 Qy 298 VTEDGLKLDTATMLQGQAKITVDKDSTVYEGSSSEIATNRLALKSOLETTSD 357
 Db 299 VTEDGLKLDTATMLQGQAKITVDKDSTVYEGSSSEIATNRLALKSOLETTSD 358
 Qy 358 FDREKLOERLAKLAGVAVKVGAPTEALKEMKLRTDNLNTRAAVEGTVAGGTL 417
 Db 359 FDREKLOERLAKLAGVAVKVGASTELKERKLRIEDALNSTRAVEGTVAGGTL 418
 Qy 418 ITVTEKVALELEGDDATGNTTIALELPYRQALNAGYGSVYDVKUNSPAGTGFN 477
 Db 419 VVNVQKVSNEAREGDIETGVNIVKLTAVPKVQIAENAGLEGSVIVERLKHAEAGVGFNG 478
 Qy 478 ATGEMDVKIGIIPVKVTRALQNLQNSASVSLTTEAVVANKPEPATPAPAMPMDP 537
 Db 479 ATNBNVWMLRQGIVDPTKVRSALQHASYVAMFLTEAUVVASIPENNEBPGM-GMM-P 536
 Qy 538 GMM 540

Db	537	Gbk	539
RESULT	13		
ABP9860	ABP9860 standard; Protein: 540 AA.		
AC	AC		
DBP9860;			
XX			
DT	24-JUL-2002 (first entry)		
DE	Staphylococcus epidermidis ORF amino acid sequence SEQ ID NO:4705.		
XX			
KW	Staphylococcus epidermidis; open reading frame; ORF; bacterial infection; antibacterial; gene therapy.		
XX			
OS	Staphylococcus epidermidis.		
PN	US6380370-B1.		
PD	30-APR-2002.		
XX			
PF	13-AUG-1998; 980US-0134001.		
XX			
PR	14-AUG-1997; 97US-055779P.		
XX			
PR	08-NOV-1997; 97US-064664P.		
PA	(EBNO-) GENOME THERAPEUTICS CORP.		
XX			
PI	Doucette-Stamm L.A., Bush D.		
XX			
PT	WPI; 2002-381255/41.		
XX			
DR	N-PSDB; ABN92405.		
XX			
PT	Novel isolated nucleic acid encoding a Staphylococcus epidermidis polypeptide, useful for diagnosing and treating bacterial infections -		
XX			
PS	DISCLOSURE; SEQ ID 4705; 267PP; English.		
XX			
CC	ABN90338 to ABN93374 represent Staphylococcus epidermidis open reading frame (ORF) nucleic acid sequences which encode the amino acid sequences given in ABP514 to ABP3790. The S. epidermidis sequences have antibacterial activity and can be used in gene therapy. The sequences can also be used in the diagnosis and treatment of bacterial infections, particularly S. epidermidis infections. The sequences can be used to screen for compounds able to interfere with the S. epidermidis life cycle or inhibit S. epidermidis infection.		
CC	N.B. The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from the USPTO web site.		
CC			
SQ	Sequence 540 AA;		
Query Match	68.9%; Score: 1813.5; DB 23; Length 540;		
Best Local Similarity	66.5%; Pred. No. 2.4e-10; Mismatches: 87; Mismatches: 93; Indels 1; Gaps 1;		
Matches	359; Conservative		
QY	1 MAKEIKFESDARQAMVRGUDMADTUVKVLGPKGRNVLEKAFGSSPLTNDGVTKETE 60		
2 MAKOLKFESDARQAMVRGUDMADTUVKVLGPKGRNVLEKAFGSSPLTNDGVTKETE 61			
Db	61 LEDFENNGAKVYEVASKNDIAGGTTATWLTQALVHEGLKVNTRGANPIGRIGIE 120		
QY	62 LEDYENNGAKLVQEVANKNTNELAGGTTATWLTQALVHEGLKVNTRGANPIGRIGIE 121		
Db	121 TATTAVERALIAQPVSKELAQVAVSSRKEVYISEANERVENDEVITIEERG 180		
QY	122 KAVOVAIRALHEISOKVENKEIAQGATSAADELGRYISEAKDVKVENDGTYIESNG 181		
Db	181 NETELEVVEVGKQDFDGVYQSYMTDNEKWDADENPILTDKVSNTQDPLIEYKL 240		
QY	182 FNTSELEVEVGKQDFDGVYQSYMTDNEKWDADENPILTDKVSNTQDPLIEYKL 241		
PT	New nucleic acid encoding heat shock protein-60 from Streptococcus, useful in vaccines, as carriers for other immunogens, as anticancer agents and for diagnosis		
PT	agents and for diagnosis		
XX			
PS	DISCLOSURE; FIG 10A-E; 176pp; English.		
XX			
CC	ABY23905-30 represent heat shock proteins (Hsps). The specification, which describes Streptococcal Hsps, designated Hsp60. These proteins, their fragments, variants and fusion proteins, are used to elicit or enhance an immune response against Streptococcus, and to elicit a similar response to a target antigen fused to the protein. Unlike other immunological targets, Hsp60 proteins are not immunosuppressive so provide an increased response to any conjugated or fused antigen. As such, where used for cancer control, they lack the side effect associated with endotoxins. They can also be used to detect specific antibodies and in treatment or prevention of tumours (e.g. sarcoma or cancers of breast, ovary, prostate, lung, pancreas or liver). The Hsp60 polynucleotide is used for recombinant production of the protein, as a source of primers and probes for detecting streptococci in standard hybridization/amplification assays, and therapeutically in gene therapy vectors.		

Match	Score	Length	DB	WPI	XX
Best Local Similarity	68.3%	Score: 1819; DB: 20; Length: 539;	DB	1999-430397/36.	XX
Matches	367; Conservative	67.7%; Pred. No. 2e-105;	Matches	New nucleic acid encoding heat shock protein-60 from <i>Streptococcus</i> , useful in vaccines, as carriers for other immunogens, as anticancer agents and for diagnosis	PT
Qy	1 MAKKEKFSDARAAMVGRGMDLADTYKVTLGPKGRNVLKAFGSPLITNDVTAKE 60	1 MAKTYLRLGEARRSKQAGVKDNLTVKVLGPKGRNVLKAFGSPLITNDVTAKE 60	Db	Disclosure: Fig 10A-E, 176pp; English.	PS
Qy	61 LEDHENNGAGLAVKPSKTDIAGDTTATLQAVHEGLKVNAGANPAGIRGIE 120	61 LEDHENNGAGLAVKPSKTDIAGDTTATLQAVHEGLKVNAGANPAGIRGIE 120	Db	AAV3905-30 represent heat shock proteins (Hsps). The specification describes <i>Streptococcal</i> Hsps, designated Hsp60. These proteins, their fragments, variants and fusion proteins, are used to elicit or enhance an immune response against <i>Streptococcus</i> , and to elicit a similar response to a target antigen fused to the protein. Unlike other immunological carriers, Hsp60 proteins are not immunosuppressive so provide an increased response to any conjugated or fused antigen. Also, where used for cancer control, they lack the side effects associated with endotoxins. They can also be used to detect specific antibodies and in treatment of tumours (e.g. sarcoma or cancers of breast, ovary, prostate, lung, pancreas or liver). The Hsp60 polynucleotide is used for recombinant production of the protein, as a source of primers and probes for detecting <i>streptococci</i> in standard hybridization/amplification assays, and therapeutically in gene therapy vectors.	CC
Qy	181 METELEVVEGKQFDGKYSQAVTNEKAVALDENPLITDKVSNIQDILPYLEVLK 240	181 METELEVVEGKQFDGKYSQAVTNEKAVALDENPLITDKVSNIQDILPYLEVLK 240	Db	CC	
Db	181 MGTELDVVEGKQFDGKYSQAVTNEKAVALDENPLITDKVSNIQDILPYLEVLK 240	181 MGTELDVVEGKQFDGKYSQAVTNEKAVALDENPLITDKVSNIQDILPYLEVLK 240	Qy	CC	
Qy	241 TNRPLLIADDVDAEPLVINKRGTGPNVVAKARGFDRKAMEDIAILGGTVT 300	241 TNRPLLIADDVDAEPLVINKRGTGPNVVAKARGFDRKAMEDIAILGGTVT 300	Db	CC	
Db	241 AGKPLLIADDIEGAMTLLVNLKQGIFTCPVGKAPGFDRKEMQDIAITGGTVS 300	241 AGKPLLIADDIEGAMTLLVNLKQGIFTCPVGKAPGFDRKEMQDIAITGGTVS 300	Qy	CC	
Qy	301 EDGLGELKDATMAGQAKITVDKDSTVVEGGSSEANRALKSQQLETTSDFR 360	301 EDGLGELKDATMAGQAKITVDKDSTVVEGGSSEANRALKSQQLETTSDFR 360	Db	CC	
Db	301 DEVGGDIEKATLDMGESEKVKVKESTIVNGRGNSEEIKNNIQKLQLEATSEFDK 360	301 DEVGGDIEKATLDMGESEKVKVKESTIVNGRGNSEEIKNNIQKLQLEATSEFDK 360	Qy	CC	
Qy	361 EKIQERIPLAKLAGGVAVKVGAPTEKPLKEMKIRTEDALMATRAVEEGIVAGGATVY 420	361 EKIQERIPLAKLAGGVAVKVGAPTEKPLKEMKIRTEDALMATRAVEEGIVAGGATVY 420	Db	CC	
Db	421 IERVVALE-GDDATGGRNVLRALEPPVRLTQVAGKPSKTDIUKNSPAGTGERAT 479	421 IERVVALE-GDDATGGRNVLRALEPPVRLTQVAGKPSKTDIUKNSPAGTGERAT 479	Qy	CC	
Db	421 INEVAKLTSBIDQEVQGINNIVLRSLEPMRQTAHAGLEGSPVIEKVKNSDAGVDA 480	421 INEVAKLTSBIDQEVQGINNIVLRSLEPMRQTAHAGLEGSPVIEKVKNSDAGVDA 480	Db	CC	
Qy	480 GEWDIMTKGIDPVKVTRSALQNAASVSLTTEAVVANKEPPATPAPMAGMDGM 539	480 GEWDIMTKGIDPVKVTRSALQNAASVSLTTEAVVANKEPPATPAPMAGMDGM 539	Db	CC	
Db	481 GEYKDMKAGIVDPTKVTRSLQNAASVSLTTEAVVADPKE- --- KEMPQAGM 539	481 GEYKDMKAGIVDPTKVTRSLQNAASVSLTTEAVVADPKE- --- KEMPQAGM 539	Qy	CC	
Qy	540 MG 541	540 MG 541	Db	CC	
Db	536 DG 537	536 DG 537	Qy	XX	
RESULT	15		Qy	Sequence 541 AA:	
ARY23910			Qy	Query Match 63.9%; Score 1702; DB: 20; Length 541;	
ID	ARY23910 standard; Protein; 541 AA.		Matches	Best Local Similarity 63.0%; Pred. No. 7.3e-102; Mismatches 113; Indels 4; Gaps 1;	
XX			Db	1 MAKKEKFSDARAAMVGRGMDLADTYKVTLGPKGRNVLKAFGSPLITNDVTAKE 60	
AC	ARY23910;		Qy	1 MAKTYLRLGEARRSKQAGVKDNLTVKVLGPKGRNVLKAFGSPLITNDVTAKE 60	
DT	22-SEP-1999 (first entry)		Db	61 LEDHENNGAGLAVKPSKTDIAGDTTATLQAVHEGLKVNAGANPAGIRGIE 120	
DE	Amino acid sequence of a heat shock protein.		Db	61 LEDPYEKGIAELVKAEVAKKDDVAGGTTTATLQAVLVRREGIYRGNVAGANPLIKRGE 120	
XX			Db	121 TATATAVRLAKIAQPVSKTEAQAQVAVSRSRKEVYSEAMERYNGDVOVIESRG 180	
KW	Heat shock protein; Hsp; immune response; immunological carrier; cancer control; tumour; sarcoma; cancer; gene therapy.		Db	121 KAVKVETELLSKSAKEVETDKQIATTAISAGDOSTGDLAEMKVNECVITSENT 180	
XX			Qy	181 METELEVVEGKQFDGKYSQAVTNEKAVALDENPLITDKVSNIQDILPYLEVLK 240	
Db			Db	181 FGLQELTEGKMDFKQYISGYFVTDKEROBAVLEDPLFILLVSSKVSTKDLPLPLERIQ 240	
Qy			Qy	241 TNRPLLIADDVDAEPLVINKRGTGPNVVAKARGFDRKAMEDIAILGGTVT 300	
Db			Db	241 AGKPLLIADDIEGAMTLLVNLKQGIFTCPVGKAPGFDRKEMQDIAITGGTVS 300	
Qy			Qy	301 EDGLGELKDATMAGQAKITVDKDSTVVEGGSSEANRALKSQQLETTSDFR 360	
Db			Db	301 EEVGLSLASDLSLUGKARVYVTTTGGAGSDATAGVATRTELENSDYOR 360	
Qy			Qy	361 EKLERIPLAKLAGGVAVKVGAPTEKPLKEMKIRTEDALMATRAVEEGIVAGGATVY 420	
Db			Db	361 EKLERIPLAKLAGGVAVKVGAPTEKPLKEMKIRTEDALMATRAVEEGIVAGGATVY 420	
Qy			Qy	421 IERVVALE-GDDATGGRNVLRALEPPVRLTQVAGKPSKTDIUKNSPAGTGERAT 480	
Db			Db	421 IPALDELKEPSEGAEVANVYVRLPLKQIAFENGGLPQVWAKYVRSNPGTGTGNGAT 480	
Qy			Qy	481 EWWNIKIGITDPVKTRSALQNAASVSLTTEAVVANKPAPATPAPMAGMDGM 540	
Db			Db	481 EYEDLKLAKIADPVKVTRSALQNAASVSLTTEAVVADPKE- --- KAPAGDPGGM 536	
PF	29-DEC-1998; 98HO-CM01203.		Qy	541 GGM 543	
XX			Db	537 GGM 539	
PR	31-DEC-1997; 97US-0001737.				
XX					
PA	(SRE-) STRESSGEN BIOTECHNOLOGIES CORP.				
PI	Mizzen L, Wisniewski J;				

Wed Apr 16 08:08:21 2003

us-09-001-737-8.rag

Page 12

Job time : 43 secs